

1. (Currently Amended.) A method for manufacturing an article comprising the steps of: a.) coating a molding surface of a mold or a mold half with a coating effective amount of a high molecular weight coating composition; b.) dispensing a monomer mixture comprising a hydrogel monomer, a silicone-containing hydrogel monomer, or a combination thereof into the mold or mold half; and c.) curing the monomer mixture and the coating composition using a dwell time of less than about 5 minutes and under conditions suitable to form an article coated with the coating composition.

2. (Original.) The method of claim 1, wherein the article is a contact lens.

3. (Currently amended). The method of claim 1 or 2, wherein the monomer mixture further comprises a at least one hydrogel monomer.

**Claim 4 (Cancelled).**

5. (Original.) The method of claim 1, wherein the molecular weight of the coating composition is greater than about 300 kD.

6. (Original.) The method of claim 1, wherein the dwell time is less than about 45 seconds.

7. (Original.) The method of claim 5, wherein the dwell time is less than about 45 seconds.

8. (Original.) The method of claim 1, wherein the coating composition further comprises a low boiling point solvent and a high boiling point solvent.

9. (Original.) The method of claim 8, wherein the coating of the molding surface is carried out by spin coating.

10. (Original.) The method of claim 9, wherein spin coating is carried out using at least about 2  $\mu$ l and no more than about 20  $\mu$ l of the coating composition.

11. (Original.) The method of claim 10, further comprising applying, subsequent to the spin coating step, a pressurized air jet to an edge of the mold.
12. (Withdrawn.) An article formed by the method of claim 1, 6, 7, or 9.
13. (Withdrawn.) A contact lens formed by the method of claim 2.
14. (Currently Amended.) A method for manufacturing an article comprising: a.) coating a molding surface of a mold or a mold half with a coating effective amount of a high molecular weight hydrophilic coating composition comprising poly(2-hydroxyethyl methacrylate); b.) dispensing a monomer mixture of comprising a hydrogel monomer, silicone-containing hydrogel monomer, or combination thereof into the mold or mold half; and c.) curing the monomer mixture and coating composition using a dwell time of less than about 5 minutes and under conditions suitable to form an article coated with the coating composition.
15. (Original.) The method of claim 14, wherein the article is a contact lens.
16. (Original.) The method of claim 15, wherein the monomer mixture comprises a hydrogel monomer.
17. (Original.) The method of claim 15, wherein the monomer mixture comprises a silicone hydrogel monomer.
18. (Original.) The method of claim 14, wherein the dwell time is less than about 45 seconds.
19. (Original.) The method of claim 17, wherein the silicone hydrogel monomer mixture comprises a reaction product of a silicone based macromer Group Transfer Polymerization product and a polymerizable mixture comprising Si<sub>8-10</sub> monomethacryloxy terminated polydimethyl siloxane, polydimethylsiloxane other than Si<sub>8-10</sub> monomethacryloxy terminated polydimethyl siloxane, and a hydrophilic monomer.

20. (Original.) The method of claim 19, wherein the silicone hydrogel monomer mixture comprises the macromer in an amount of about 15 to about 25 wt percent, the Si<sub>8-10</sub> monomethacryloxy terminated polydimethyl siloxane in an amount of about 20 to about 30 wt percent; methacryloxypropyl tris(trimethyl siloxy) silane in an amount of about 15 to about 25 wt percent; N,N-dimethyl acrylamide in an amount of about 20 to about 30 wt percent; 2-hydroxy ethyl methacrylate in an amount of about 2 to about 7 wt percent; tetraethyleneglycol dimethacrylate in an amount of about 0 to about 5 wt percent and poly(N-vinyl pyrrolidinone) in an amount of about 0 to about 5 weight percent.

21. (Original.) The method of claim 14, 15, 16, 17, 18, 19, or 20 wherein the coating composition comprises poly(vinyl alcohol), polyethylene oxide, poly(2-hydroxyethyl methacrylate), poly(acrylic acid), poly(methacrylic acid), poly(maleic acid), poly(itaconic acid), poly(acrylamide), poly(dimethylacrylamide), carboxymethylated polymers, polystyrene sulfonic acid, polysulfonate polymers, polysaccharides, glucosamine glycans, block or random copolymers thereof, or mixtures thereof.

22. (Cancelled.)

23. (Original.) The method of claim 15, wherein the coating composition further comprises a low boiling point solvent and a high boiling point solvent.

24. (Original.) The method of claim 23, wherein the coating of the molding surface is carried out by spin coating.

25. (Original.) The method of claim 24, wherein spin coating is carried out using at least about 2  $\mu$ l and no more than about 20  $\mu$ l of the coating composition.

26. (Original.) The method of claim 25, further comprising applying, subsequent to the spin coating step, a pressurized air jet to an edge of the mold.

27. (Withdrawn.) An article formed by the method of claim 14.

28. (Withdrawn.) A contact lens formed by the method of claim 15, 18, 19, or 20.
29. (Withdrawn.) A contact lens formed by the method of claim 21.
30. (Withdrawn.) A contact lens formed by the method of claim 22.
31. (Currently Amended.) A method for manufacturing contact lenses comprising: a.) coating a molding surface of a mold or a mold half with a coating effective amount of a hydrophilic coating composition having a molecular weight of greater than about 300 k $\text{\u0333}$ ; b.) dispensing a mixture comprising a hydrogel monomer, a silicone-containing hydrogel monomer, or a combination thereof into the mold or mold half; and c.) curing the monomer mixture and coating composition using a dwell time of less than about 45 seconds and under conditions suitable to form a contact lens coated with the coating composition, wherein the formed lens exhibits physiological compatibility.
32. (Currently Amended.) The method of claim 31, wherein the monomer mixture further comprises at least one hydrogel monomer.
33. (Cancelled.)
34. (Original.) The method of claim 33, wherein the silicone hydrogel monomer mixture comprises a reaction product of a silicone based macromer Group Transfer Polymerization product and a polymerizable mixture comprising Si<sub>8-10</sub> monomethacryloxy terminated polydimethyl siloxane, polydimethylsiloxane other than Si<sub>8-10</sub> monomethacryloxy terminated polydimethyl siloxane, and a hydrophilic monomer.
35. (Original.) The method of claim 34, wherein the silicone hydrogel monomer mixture comprises the macromer an amount of about 15 to about 25 wt percent, the Si<sub>8-10</sub> monomethacryloxy terminated polydimethyl siloxane in an amount of about 20 to about 30 wt percent; methacryloxypropyl tris(trimethyl siloxy) silane in an amount of about 15 to about 25 wt percent; N,N-dimethyl acrylamide in an amount of about 20 to about 30 wt percent; 2-hydroxy ethyl methacrylate in an amount of about 2 to about 7 wt percent;

tetraethyleneglycol dimethacrylate in an amount of about 0 to about 5 wt percent and poly(N-vinyl pyrrolidinone) in an amount of about 0 to about 5 weight percent.

36. (Currently Amended.) The method of claim 31, 32, 33, 34, or 35, wherein the coating composition comprises poly(vinyl alcohol), polyethylene oxide, poly(2-hydroxyethyl methacrylate), poly(acrylic acid), poly(methacrylic acid), poly(maleic acid), poly(itaconic acid), poly(acrylamide), poly(dimethacrylamide), carboxymethylated polymers, polystyrene sulfonic acid, polysulfonate polymers, polysaccharides, glucose amino glycans, block or random copolymers thereof, or mixtures thereof.

37. (Original.) The method of claim 36, wherein the coating composition comprises poly(2-hydroxyethyl methacrylate).

38. (Original.) The method of claim 36, wherein the coating composition further comprises a low boiling point solvent and a high boiling point solvent.

39. (Original.) The method of claim 38, wherein the coating of the molding surface is carried out by spin coating.

40. (Original.) The method of claim 39, wherein spin coating is carried out using at least about 2  $\mu$ l and no more than about 20  $\mu$ l of the coating composition.

41. (Original.) The method of claim 40, further comprising applying, subsequent to the spin coating step, a pressurized air jet to an edge of the mold.

42. (Withdrawn.) A contact lens formed by the method of claim 31, 32, 33, 34, or 35.

43. (Withdrawn.) A contact lens formed by the method of claim 36.

44. (Withdrawn.) A contact lens formed by the method of claim 37.

45. (Withdrawn.) A contact lens formed by the method of claim 38.

46. (Withdrawn.) A contact lens formed by the method of claim 39.

47. (Withdrawn.) The contact lens of claim 42, wherein the coating composition comprises a coating of a dry film thickness of not less than about 5 nm and not more than about 70 nm.

48. (Withdrawn.) The contact lens of claim 43, wherein the coating composition comprises a coating of a dry film thickness of not less than about 5 nm and not more than about 70 nm.

49. (Withdrawn.) The contact lens of claim 44, wherein the coating composition comprises a coating of a dry film thickness of not less than about 5 nm and not more than about 70 nm.

50. (Withdrawn.) The contact lens of claim 45, wherein the coating composition comprises a coating of a dry film thickness of not less than about 5 nm and not more than about 70 nm.

51. (Withdrawn.) The contact lens of claim 4, wherein the coating composition comprises a coating of a dry film thickness of not less than about 5 nm and not more than about 70 nm.

52. (Currently Amended.) The method of claim 1 wherein the coating composition has a Brookfield viscosity of about 17.7 cP at 25°C.

53. (Original.) The method of claim 1 wherein the coating composition has a viscosity of at least about 4 cP at 25°C.

54. (Original.) The method of claim 1 wherein the coating composition has a viscosity of greater than about 1 cP at 25°C.

55. (Original.) The method of claim 8 wherein the low boiling point solvent comprises ethanol and the high boiling point solvent comprises ethyl lactate.

56. (Original.) The method of claim 8 wherein the low boiling point solvent has a boiling point of less than about 90°C.

57. (Original.) The method of claim 8 wherein the low boiling point solvent comprises ethanol.

58. (Original.) The method of claim 8 wherein the low boiling point solvent is selected from the group consisting of n-methyl pyrrolidone, acetone, chloroform, methanol, ethanol, isopropanol, tert-butanol and combinations thereof.

59. (Original.) The method of claim 8 wherein the high boiling point has a boiling point of greater than about 100°C.

60. (Original.) The method of claim 8 wherein the high boiling point solvent comprises ethyl lactate.

61. (Original.) The method of claim 8 wherein the high boiling point solvent is selected from the group consisting of methyl lactate, ethyl lactate, isopropyl lactate, ethylene glycol, polyethylene glycol, propylene glycol, dimethyl formamide, tetrahydrogeraniol, 1-butanol, 1-pentanol, 1-hexanol, 1-octanol, 3-methyl-3-pentanol, dimethyl-3-octanol, 3-methoxy-1-butanol, 1,2-butanediol, 1,4-butanediol, 1,3-hexanediol, water and combinations thereof.

62. (Original.) The method of claim 8 wherein the low boiling point solvent and the high boiling point solvent are present at a ratio of about 1:1.

63. (Original.) The method of claim 55 wherein the low boiling point solvent and the high boiling point solvent are present at a ratio of about 1:1.

64. (Original.) The method of claim 14 wherein the coating composition has a viscosity of about 17.7 cP at 25°C.

65. (Original.) The method of claim 14 wherein the coating composition has a viscosity of at least about 4 cP at 25°C.

66. (Original.) The method of claim 14 wherein the coating composition has a viscosity of greater than about 1 cP at 25°C.
67. (Original.) The method of claim 23 wherein the low boiling point solvent comprises ethanol and the high boiling point solvent comprises ethyl lactate.
68. (Original.) The method of claim 23 wherein the low boiling point solvent has a boiling point of less than about 90°C.
69. (Original.) The method of claim 23 wherein the low boiling point solvent comprises ethanol.
70. (Original.) The method of claim 23 wherein the low boiling point solvent is selected from the group consisting of n-methyl pyrrolidone, acetone, chloroform, methanol, ethanol, isopropanol, tert-butanol and combinations thereof.
71. (Original.) The method of claim 23 wherein the high boiling point has a boiling point of greater than about 100°C.
72. (Original.) The method of claim 23 wherein the high boiling point solvent comprises ethyl lactate.
73. (Original.) The method of claim 23 wherein the high boiling point solvent is selected from the group consisting of methyl lactate, ethyl lactate, isopropyl lactate, ethylene glycol, polyethylene glycol, propylene glycol, dimethyl formamide, tetrahydrogeraniol, 1-butanol, 1-pentanol, 1-hexanol, 1-octanol, 3-methyl-3-pentanol, dimethyl-3-octanol, 3-methoxy-1-butanol, 1,2-butanediol, 1,4-butanediol, 1,3-hexanediol, water and combinations thereof.
74. (Original.) The method of claim 23 wherein the low boiling point solvent and the high boiling point solvent are present at a ratio of about 1:1.
75. (Original.) The method of claim 65 wherein the low boiling point solvent and the high boiling point solvent are present at a ratio of about 1:1.

76. (Original.) The method of claim 31 wherein the coating composition has a viscosity of about 17.7 cP at 25°C.
77. (Original.) The method of claim 31 wherein the coating composition has a viscosity of at least about 4 cP at 25°C.
78. (Original.) The method of claim 31 wherein the coating composition has a viscosity of greater than about 1 cP at 25°C.
79. (Original.) The method of claim 38 wherein the low boiling point solvent comprises ethanol and the high boiling point solvent comprises ethyl lactate.
80. (Original.) The method of claim 38 wherein the low boiling point solvent has a boiling point of less than about 90°C.
81. (Original.) The method of claim 38 wherein the low boiling point solvent comprises ethanol.
82. (Original.) The method of claim 38 wherein the low boiling point solvent is selected from the group consisting of n-methyl pyrrolidone, acetone, chloroform, methanol, ethanol, isopropanol, tert-butanol and combinations thereof.
83. (Original.) The method of claim 38 wherein the high boiling point has a boiling point of greater than about 100°C.
84. (Original.) The method of claim 38 wherein the high boiling point solvent comprises ethyl lactate.
85. (Original.) The method of claim 38 wherein the high boiling point solvent is selected from the group consisting of methyl lactate, ethyl lactate, isopropyl lactate, ethylene glycol, polyethylene glycol, propylene glycol, dimethyl formamide, tetrahydrogeraniol, 1-butanol, 1-pentanol, 1-hexanol, 1-octanol, 3-methyl-3-pentanol, dimethyl-3-octanol, 3-

methoxy-1-butanol, 1,2-butanediol, 1,4-butanediol, 1,3-hexanediol, water and combinations thereof.

86. (Original.) The method of claim 38 wherein the low boiling point solvent and the ~~high~~ boiling point solvent are present at a ratio of about 1:1.

87. (Original.) The method of claim 75 wherein the low boiling point solvent and the ~~high~~ boiling point solvent are present at a ratio of about 1:1.